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FIGURES

Fig. 1.

A

TatA (Eco)	M- SCISITNCILITDAVAVVWVHPCT KKLG-----	26
TatE (Eco)	M- GEISITNCILITDAVAVVWVHPCT KKLR-----	26
TatAy (Bsu)	M- PIGPQCSLAVVAVVWVHPCT KKLP-----	25
TatAd (Bsu)	MFS SLIGIPQCSLAVVAVVWVHPCT KKLP-----	27
TatAc (Bsu)	M- SLISITNCILITDAVAVVWVHPCT DKLP-----	25
TatB (Eco)	M- PIGPQCSLAVVAVVWVHPCT QRLPVAVKTVAGWIRALRSLATTVQNELTQELKLO	49
	* *	
TatA (Eco)	-----SIGSDLGASIKGFKKAMSDDE-----PKQDKTSQDADFTAKTI	64
TatE (Eco)	-----TLGGDLGAAIKGFKKAMNDDD-----A-AAKKGADVLDQAEKL	63
TatAy (Bsu)	-----ELGKAAGDTLREFKNATKGLT-----SDEEEKKEDQ-----	57
TatAd (Bsu)	-----EIGRAAKRTLLEFKSATKSLV-----SGDEKEEKSaelTAVK-	64
TatAc (Bsu)	-----ALGRAAGKALSEFKQATSGLT-----QDIRKNSEN-----K-	57
TatB (Eco)	EFQDSLKKVEKASLTNLTPELKASMDLRQAESMKRSYVANDPEKASDEAHTIHNP	114
 *	
TatA (Eco)	ADKQADTNQE-----QAKTEDAKRHDKEQV	89
TatE (Eco)	SHKE-----	67
TatAy (Bsu)	-----	57
TatAd (Bsu)	-----QDKNAG	70
TatAc (Bsu)	-----EDKQM-	62
TatB (Eco)	VVKDNEAAHEGVTFAAAQTQASSPEQKPEPTTPEPVVKPAADAEPKTAAPSPSSSDKP	171

B

TatC (Eco)	MSVEDTQ--PLITHLIELRK RIENCI LAVVWVHPCT NDIYH -LVSAPLIK	51
TatCy (Bsu)	MTRMKVNQMSLLEHIAELRK RIENCI LAVVWVHPCT KPII VYLQETDEAK	50
TatCd (Bsu)	MDKKETH---LIGHLEELRR RIENCI LAVVWVHPCT QDIYD WLIRDLDGK	51
	* *	
TatC (Eco)	QLPQGSTMIATDVASPFFTP IKLTF IVSV SLIS APV LYQV AFIAPALYKHERR	105
TatCy (Bsu)	QL---TLNAFNLTD PIAVEMO PA THIG IV MSPM YQLWAFVSPGLYEKERK	104
TatCd (Bsu)	-----LAVLGPSE PIAVEMO PA THIG IV MSPM YQLWRFVAPALTKTERK	98
 *	
TatC (Eco)	LVPPLLV---SSSL PIAVEMO PA THIG IV MSPM YQLWRFVAPALTKTERK	155
TatCy (Bsu)	VTLSYI---P SLIS APV LYQV AFIAPALYKHERR	155
TatCd (Bsu)	VTIMYIYIP SLIS APV LYQV AFIAPALYKHERR	151
	* *	
TatC (Eco)	SPVMAH EMARGVSE PIAVEMO PA THIG IV MSPM YQLWRFVAPALTKTERK	209
TatCy (Bsu)	HFL PIAVEMO PA THIG IV MSPM YQLWRFVAPALTKTERK	209
TatCd (Bsu)	RFM PIAVEMO PA THIG IV MSPM YQLWRFVAPALTKTERK	205
	* *	
TatC (Eco)	PDV ESQ TEAD PM QCM IGV TR SG Y GKGRNREEENDAEAESEKTEE	258
TatCy (Bsu)	PDV ESQ TEAD PM QCM IGV TR SG Y AYRKAQKSSAADRDVSSG-----Q	254
TatCd (Bsu)	PDV ESQ TEAD PM QCM IGV TR SG Y YKKRMRE-----ETAAA-----A	245
	* *	

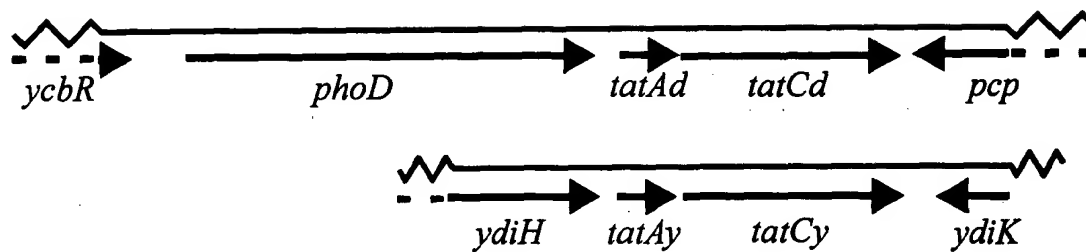
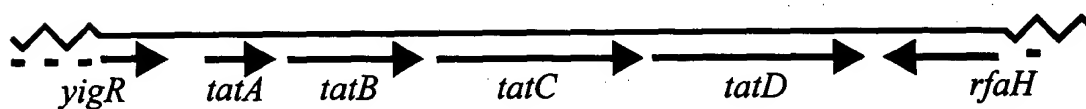
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Fig. 2.

A *B. subtilis*B *E. coli*

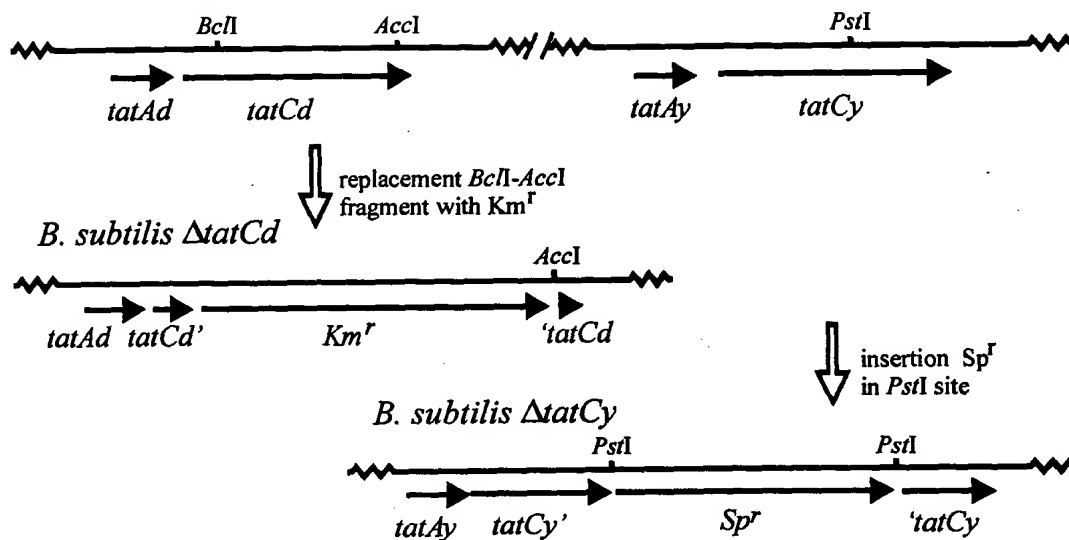
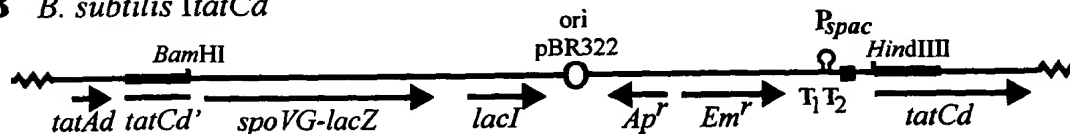
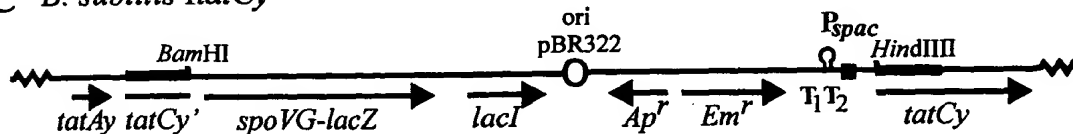
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Fig. 3.

A *B. subtilis* 168B *B. subtilis* 1*tatCd*C *B. subtilis* 1*tatCy*

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Fig. 4.

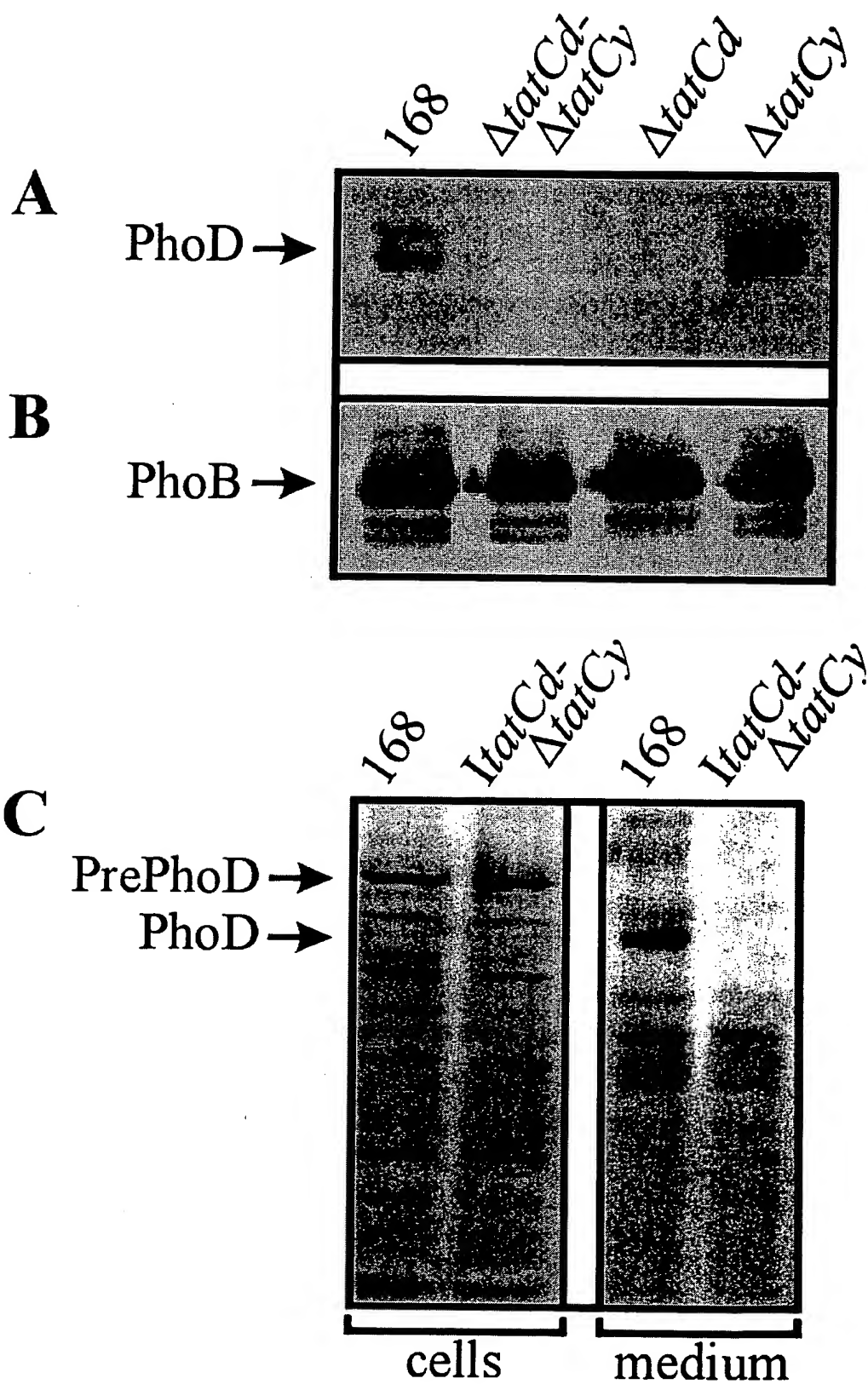
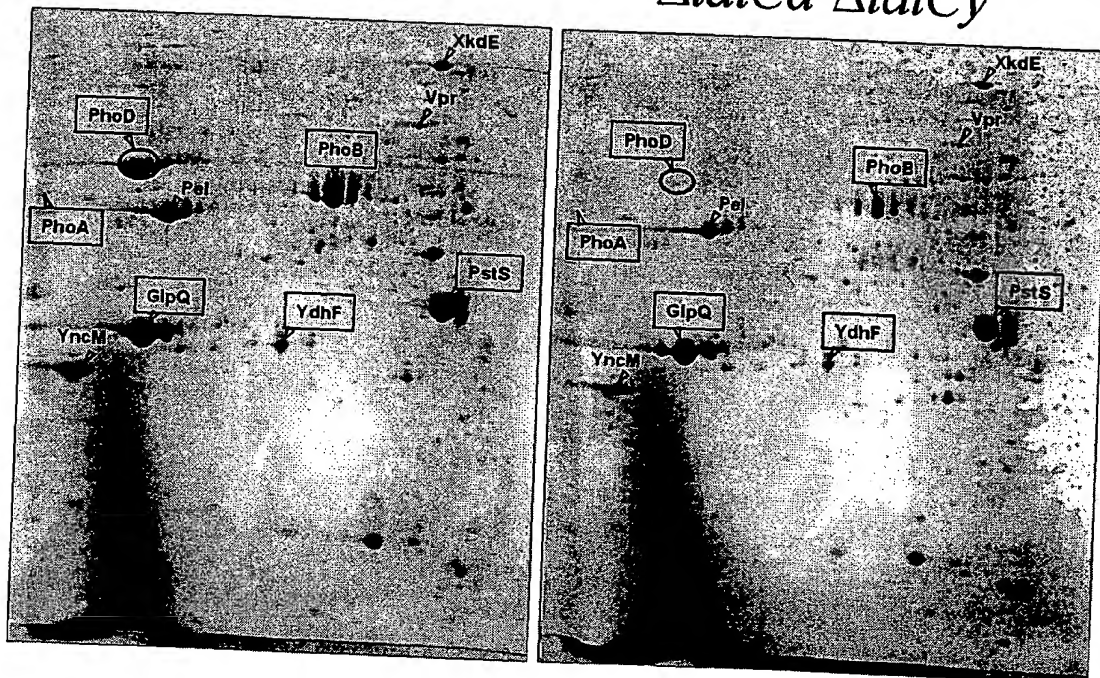
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Fig. 5.

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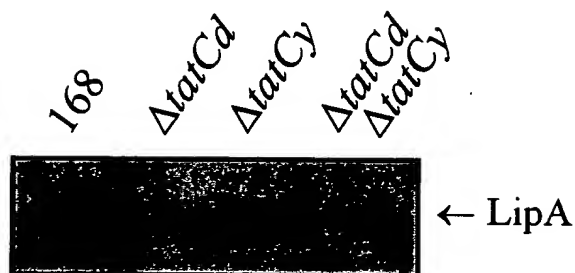
$\Delta tatCd-\Delta tatCy$



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FIGURE 6



Tat-dependent secretion of the *B. subtilis* lipase LipA. *B. subtilis* 168 (parental strain), *B. subtilis* $\Delta tatCd$, *B. subtilis* $\Delta tatCy$, or *B. subtilis* $\Delta tatCd \Delta tatCy$ were grown in TY-medium to end-exponential growth phase. To study the secretion of LipA, *B. subtilis* cells were separated from the growth medium by centrifugation. Proteins in the growth medium were concentrated 20-fold upon precipitation with trichloroacetic acid, and samples for polyacrylamide gel electrophoresis (SDS-PAGE) were prepared. Secreted LipA in the growth medium was visualized by SDS-PAGE and Western blotting, using LipA-specific antibodies.

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FIGURE 7

Predicted twin-arginine (RR-)signal peptides of *B. subtilis*¹

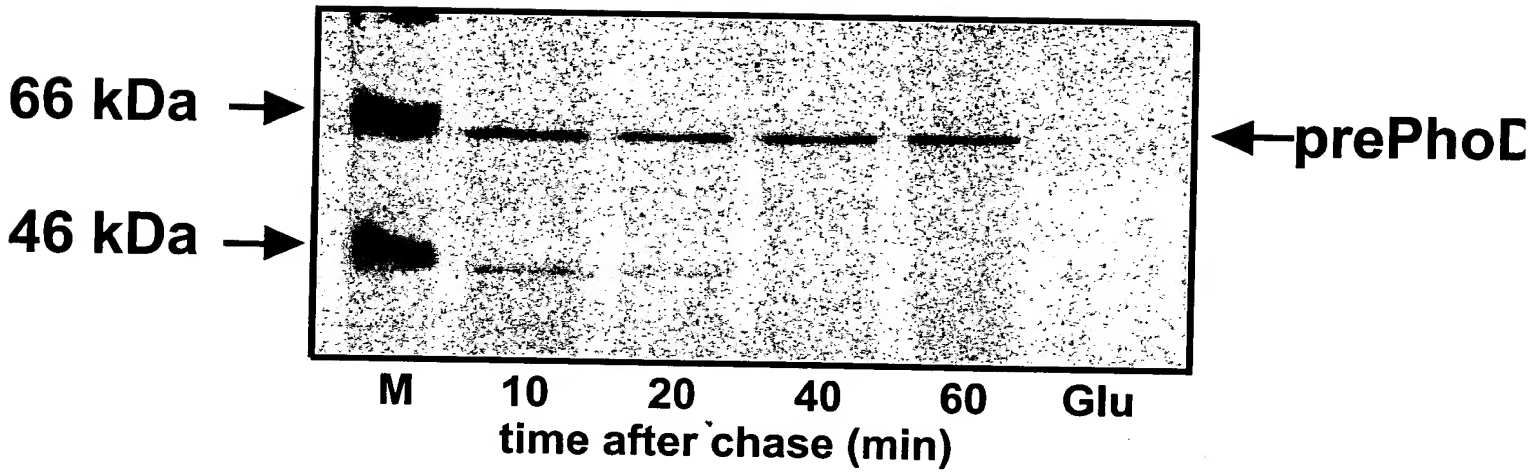
Protein	N	h	RR-Motif	H	h	C
AlbB	1	0.1	RRILL	27	2.0	AIA
AmyX TM	9	-0.8	RRSFE	15	1.1	-
AppB TM	8	0.5	RRTLm	19	2.3	-
LipA	7	-1.1	RRIIA	19	1.2	AKA
OppB TM	8	-0.6	RRLVY	24	2.0	-
PbpX	2	-2.2	RRRKL	14	2.9	WNA
PhoD	3	-1.3	RRKFI	17	0.9	VGA
QcrA TM	1	-1.1	RRQFL	19	1.3	-
TlpA TM	1	-0.8	RRLII	21	2.4	-
WapA ^W	1	-3.0	RRNFK	18	2.3	VLA
WprA	8	-1.7	RRKFS	20	1.9	AAA
YceA TM	1	-0.4	RR AFL	21	2.2	-
YesM TM	1	-1.5	RRMKI	20	2.4	QYA
YesW	1	-1.3	RRSCL	19	2.0	VKA
YfkN TM	1	-1.2	RRTHV	17	1.7	IHA
YkpC	8	-1.0	RRVAI	17	2.3	SLA
YkuE	1	-1.3	RRQFL	17	1.0	GYA
YmaC	7	0.0	RRFLL	15	2.4	YSL
YubF TM	9	-2.7	RRNTV	23	2.0	-
YuiC	8	0.2	RRLLM	20	1.9	IEA
YvhJ TM	2	-1.7	RRKIL	18	2.5	-
YwbN	1	-1.8	RRDIL	23	1.4	QTA

¹ The listed signal peptides contain, in addition to the twin-arginines, at least one other residue of the consensus sequence (R-R-X- ϕ - ϕ ; printed in bold). The number of residues in the N- and H-domains of each signal peptide, and the average hydrophobicity (h) of each of these domains, as determined by the algorithms of Kyte and Doolittle (Kyte, J., and R. F. Doolittle [1982] A simple method for displaying the hydropathic character of a protein. J. Mol. Biol. 157:105-32), are indicated. Furthermore, the RR-motifs in the N-domain, and SPase I recognition sites in the C-domain (*ie.* positions -3 to -1 relative to the predicted SPase cleavage site) are shown. Proteins lacking a (putative) SPase I cleavage site, some of which contain additional transmembrane domains, are indicated with "TM". One protein containing cell wall binding repeats is indicated with "W".

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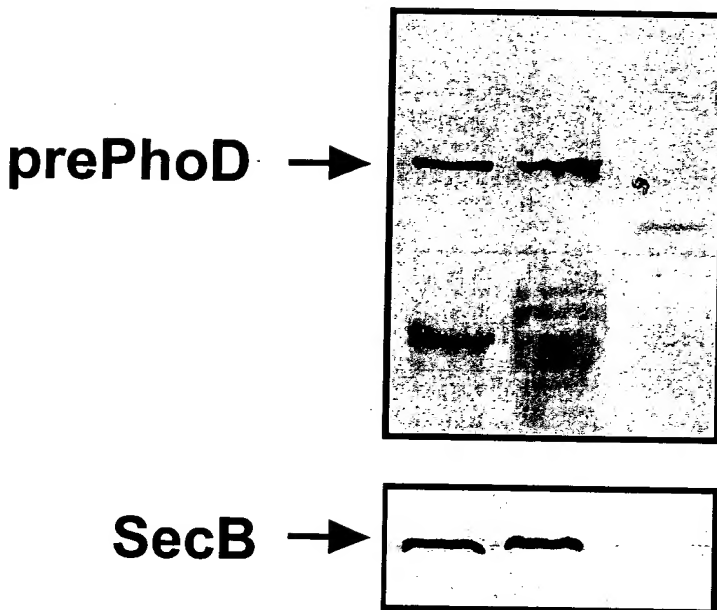
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A



B

TOLEDO 2E245660



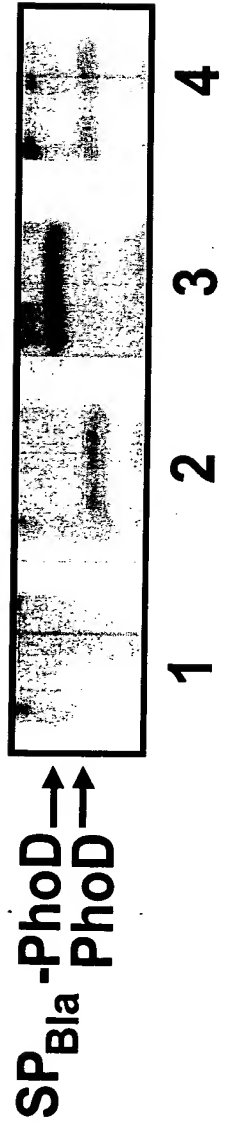
Proteinase K	-	+	+
Triton X-100	-	-	+

Figure **8**

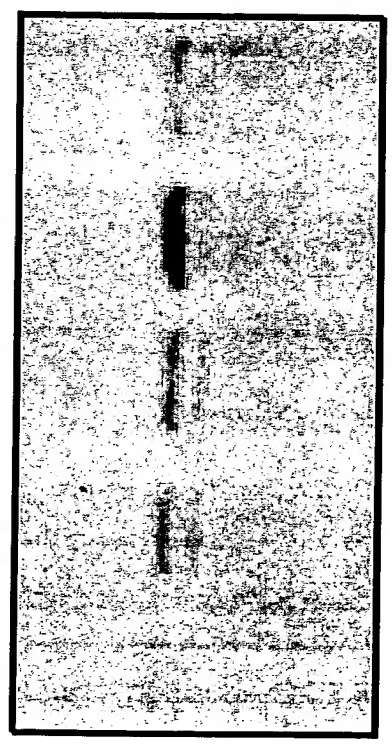
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TOPT60" 2E245660

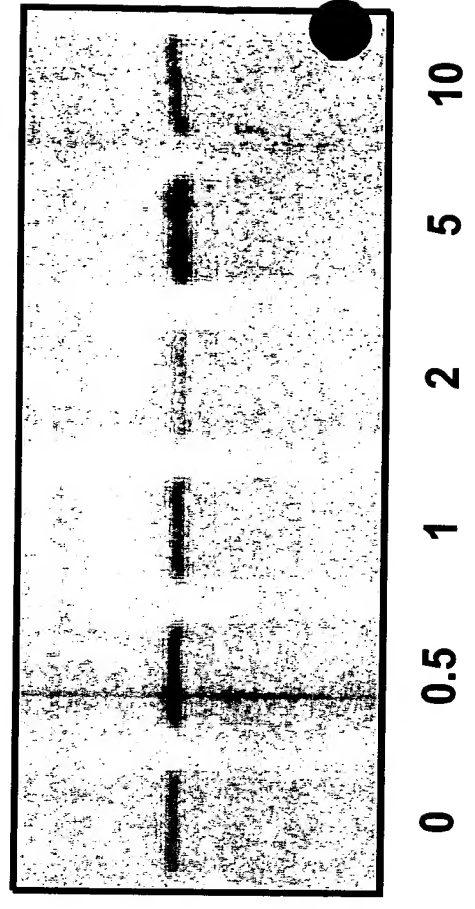
A



B untreated



C + NaN₃



time after chase

Figure 9

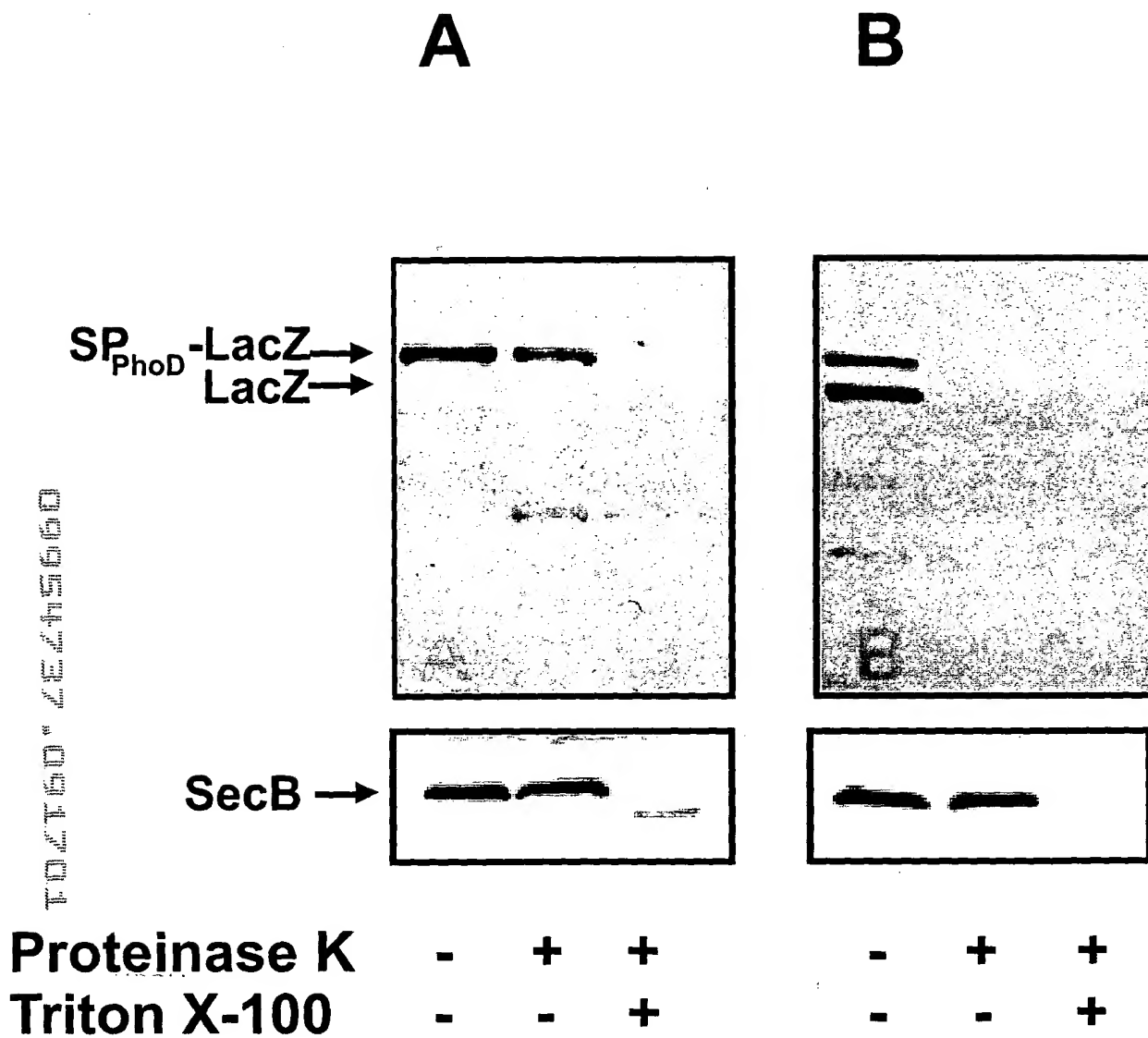


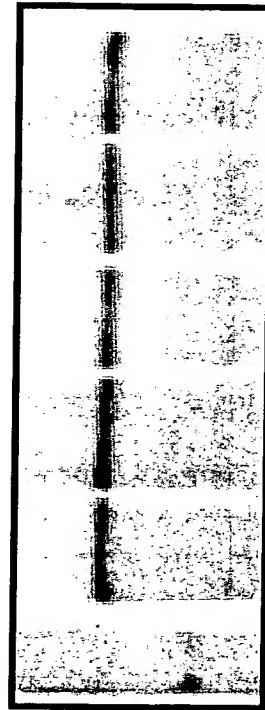
Figure 10

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FO2760" ZE245660

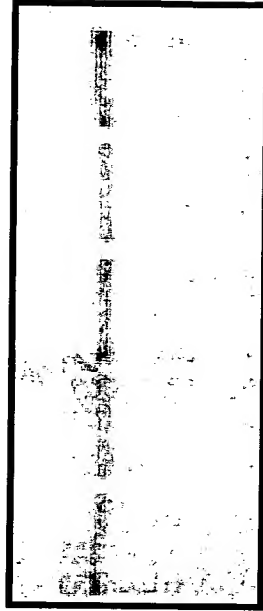
A



97.4 kDa →

M 0 2 5 10 20
time after chase (min)

B



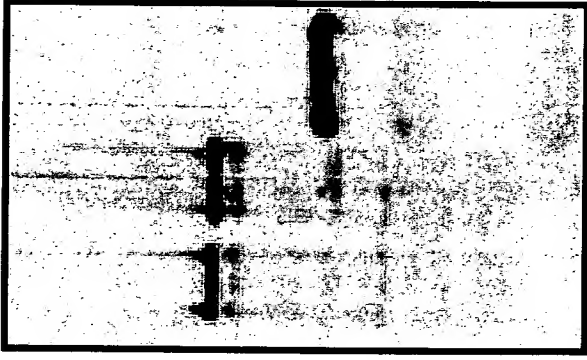
SP^{PhoD}-LacZ
←

0 2 5 10 20

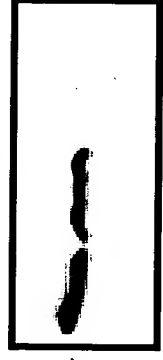
Figure 11

A
+nigericin

SP_{PhoD}-LacZ
LacZ



B
+NaN₃



SecB



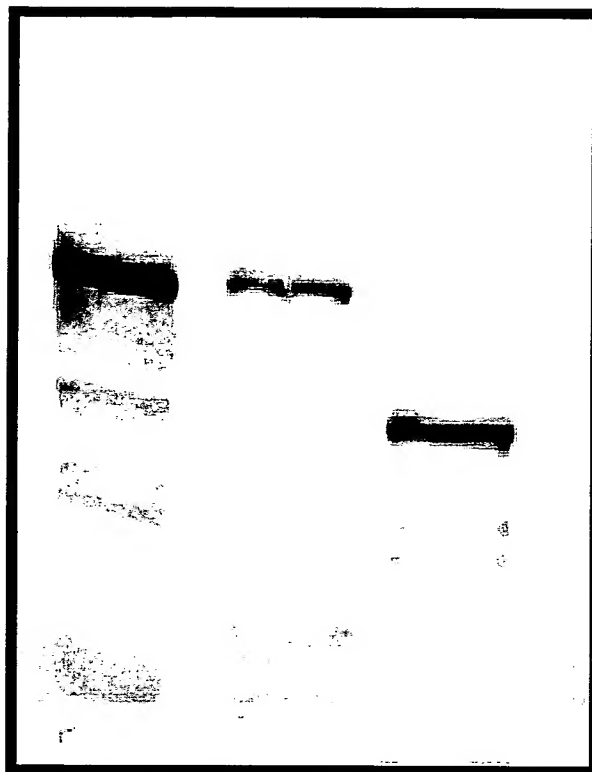
Proteinase K	-	+	+	+	-	+
Triton X-100	-	-	-	+	-	+

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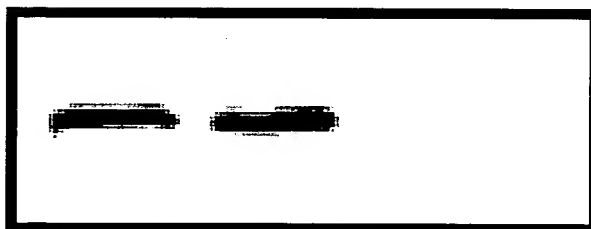
Figure 12

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SP_{PhoD}-LacZ →



SecB →



Proteinase K
Triton X-100

-	+	+
-	-	+

Figure 13

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Figure 14
Homologs in *B. alcalophilus*

TatA

MGGLSVGSVVLIALVALLIFGPKKLPELGKAAGSTLREFKNATK
GLADDDDDTKSTNVQKEKA

TatC

MTMMTPNQQTSKKKKRKGRKGRVPMQDMSIMDHAEELRRRIF
VVLAFFIVALIGGFFLAVPVITFLQNSPQAADMPFNAFRLTDPLRV
YMNFAVITALVLIIPVILYQLWAFVSPGLKENEQKATLAYIPIAFL
LFLAGIAFSYFILLPFVISFMGQMADRLEINEMYGINEYFSFLFQL
TIPFGLLFQLPVVVMFLTRLGVVTPPTFLRKIRKYAYFALLVIAGII
TPPELTSHLFVTVPMLILYEISITISAITYRKYHGTTHNGQESAK

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